

INTERNATIONAL SEARCH REPORT

International Application No
PCT/DE2005/000550

A. CLASSIFICATION OF SUBJECT MATTER IPC 7 C12Q1/68		
According to International Patent Classification (IPC) or to both national classification and IPC		
B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) IPC 7 C12Q		
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched		
Electronic data base consulted during the International search (name of data base and, where practical, search terms used) EPO-Internal, WPI Data, PAJ, EMBASE, BIOSIS, Sequence Search		
C. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X A	<p>WO 02/44426 A (REGENTS OF THE UNIVERSITY OF MICHIGAN; THE UNIVERSITY OF CHICAGO; NUNE) 6 June 2002 (2002-06-06) page 62, line 21 – page 65 page 119 page 125 claims 1-12; figures 11,17,23,25,26; examples 9,10; tables 1-6; sequences 33,56,58 & DATABASE Geneseq 'Online! 16 October 2002 (2002-10-16), "Nod2 exon 11 DNA sequence SEQ ID No 105." retrieved from EBI accession no. GSN:ABT05811 Database accession no. ABT05811</p> <p>-----</p> <p>-/-</p>	1-4,6-8 5
<input checked="" type="checkbox"/> Further documents are listed in the continuation of box C.		<input checked="" type="checkbox"/> Patent family members are listed in annex.
* Special categories of cited documents :		
A document defining the general state of the art which is not considered to be of particular relevance *E* earlier document but published on or after the International filing date *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) *O* document referring to an oral disclosure, use, exhibition or other means *P* document published prior to the International filing date but later than the priority date claimed		
T later document published after the International filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention *X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone *Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art. *&* document member of the same patent family		
Date of the actual completion of the International search	Date of mailing of the International search report	
17 August 2005	25/08/2005	
Name and mailing address of the ISA European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax. (+31-70) 340-3016	Authorized officer Schmitt, A	

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C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 2004/053263 A1 (ABREU MARIA T ET AL) 18 March 2004 (2004-03-18) claims 1-15; figures 5-7; example 2; table 3; sequences 48,50,52 -----	1-4,6-8
A	claims 1-15; figures 5-7; example 2; table 3; sequences 48,50,52	5
X	LESAGE SUZANNE ET AL: "CARD15/NOD2 mutational analysis and genotype-phenotype correlation in 612 patients with inflammatory bowel disease." AMERICAN JOURNAL OF HUMAN GENETICS. APR 2002, vol. 70, no. 4, April 2002 (2002-04), pages 845-857, XP002340892 ISSN: 0002-9297 the whole document -----	1-4,6-8
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X	HAMPE J ET AL: "Association of NOD2 (CARD 15) genotype with clinical course of Crohn's disease: a cohort study" LANCET THE, LANCET LIMITED. LONDON, GB, vol. 359, no. 9318, 11 May 2002 (2002-05-11), pages 1661-1665, XP004790813 ISSN: 0140-6736 the whole document -----	6-8
X	RAHMAN P ET AL: "CARD15: a pleiotropic autoimmune gene that confers susceptibility to psoriatic arthritis." AMERICAN JOURNAL OF HUMAN GENETICS. SEP 2003, vol. 73, no. 3, September 2003 (2003-09), pages 677-681, XP002340893 ISSN: 0002-9297 the whole document -----	1-4,6-8
X	HUGOT JEAN-PIERRE ET AL: "Association of NOD2 leucine-rich repeat variants with susceptibility to Crohn's disease" NATURE, MACMILLAN JOURNALS LTD. LONDON, GB, vol. 411, no. 6837, 2001, pages 599-603, XP002177308 ISSN: 0028-0836 cited in the application the whole document -----	1-4,6-8
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X	OGURA YASUNORI ET AL: "A frameshift mutation in NOD2 associated with susceptibility to Crohn's disease" NATURE, MACMILLAN JOURNALS LTD. LONDON, GB, vol. 411, no. 6837, 2001, pages 603-606, XP002177309 ISSN: 0028-0836 cited in the application the whole document -----	1-4, 6-8
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Information on patent family members

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